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REQUEST FOR FILING APPLICATION

Under Rule 53(a), (b) & (f)

(No Filing Fee or Oath/Declaration)

(Do NOT use for Provisional or PCT Applications)

Use for Design or Utility Applications

PATENT
APPLICATION

RULE 53(f) NO DECLARATION

Assistant Commissioner of Patents

Atty. Dkt.

PM 265189

9143US/CON/
WO

M#

Client Ref

and Trademarks
Washington, DC 20231

Date:

December 10, 1999

Sir:

1. This is a Request for filing a new Patent Application(☐ Design ☒ Utility) entitled:

2. (Complete) Title: PROCESS FOR THE PREPARATION OF AMPICILLIN

without a filing fee or Oath/Declaration but for which is enclosed the following:

3. ☒ Abstract 2 page(s).4. 13 Pages of Specification (only spec. and claims); 5. ☐ Specification in non-English language

6. 10 Numbered claim(s); and

7. ☐ 0 sheet(s) per set; ☐ 1 set informal; 8. ☐ formal of size: ☐ A4 ☐ 11"

Drawings:

DOMESTIC/INTERNATIONAL priority is claimed under 35 USC 119(e)/120/365(c) based on the following provisional, nonprovisional and/or PCT international application(s):

Application No.	Filing Date	Application No.	Filing Date
(1) PCT/NL98/00295	May 25, 1998	(2)	
(3)		(4)	
(5)		(6)	

0. FOREIGN priority is claimed under 35 USC 119(a)-(d)/365(b) based on filing in The Netherlands

Application No.	Filing Date	Application No.	Filing Date
(1) 1006266	June 10, 1997	(2)	
(3)		(4)	
(5)		(6)	

11. (No.) Certified copy (copies): ☐ attached; ☐ previously filed (date) filed on12. ☐ This is a reissue of Patent No. _____13. ☐ See top first page re prior Provisional, National, International application(s) (X box only if info is there and do not complete corresponding item 14 or 15.)14. ☒ Amend the specification by inserting before the first line -- This is a ☐ Continuation-in-Part
☐ Divisional ☒ Continuation ☐ Substitute Application (MPEP 201.09) of:14(a) ☐ National Appln. No. / filed -- (M#)
14(b) ☒ International Appln. No. PCT/NL98/00295 filed May 25, 199815. ☐ Amend the specification by inserting before the first line: --This application claims the benefit of U.S. Provisional Application No. 60/ , filed --16. Extension to date: ☐ concurrently filed ☐ not needed ☐ previously filed

17. ☐ Prior application is assigned to

by Assignment recorded _____ Reel _____ Frame _____

18. ☒ **Attached: Form PTO-1449, a copy of ISR and International Preliminary Examination Report.**

19. This application is made by the following named inventor(s) (Double check instructions for accuracy.):

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20. NOTE: FOR ADDITIONAL INVENTORS, check box ☐ and attach sheet with same information regarding additional inventors.

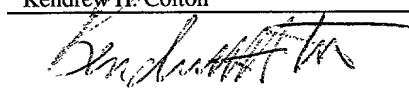
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NOTE: File in duplicate with 2 post card receipts (PAT-103) & attachments

APPLICATION UNDER UNITED STATES PATENT LAWS

Atty. Dkt. No. PM 265189
(M#)

Invention: **PROCESS FOR THE PREPARATION OF AMPICILLIN**

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This is a:

- ☐ Provisional Application
- ☐ Regular Utility Application
- ☒ Continuing Application
- ☐ PCT National Phase Application
- ☐ Design Application
- ☐ Reissue Application
- ☐ Plant Application
- ☐ Substitute Specification
Sub. Spec. filed _____
in App. No. _____ / _____
- ☐ Marked Up Specification re
Sub. Spec. filed _____
in App. No. _____ / _____

SPECIFICATION

5 PROCESS FOR THE PREPARATION OF AMPICILLIN

The invention relates to a process for the preparation of ampicillin in which 6-aminopenicillanic acid (6-APA) is subjected to an enzymatic acylation reaction with the aid of a phenylglycine derivative, with the total concentration of the 6-APA present in the reaction mixture, plus ampicillin, being greater than 250 mM, the concentration of 6-APA in solution being kept lower than 300 mM and the molar ratio of acylation agent to 6-APA which is employed being less than 2.5.

WO-A-92/01061 describes the preparation of β -lactam derivatives, including ampicillin, via enzymatic acylation of a β -lactam nucleus, for example 6-APA, at high concentrations of acylation agent plus β -lactam derivative. The concentration of the β -lactam nucleus is kept relatively low. From the examples it can be deduced that high conversions are achieved at a high molar ratio of acylation agent to β -lactam nucleus, whereas the conversion is significantly lower at a lower molar ratio of acylation agent to β -lactam nucleus. A disadvantage of the use of a high molar ratio of acylation agent to β -lactam nucleus is that large amounts of acylation agent are lost because of hydrolysis of the acylation agent. In addition it has been found that upgrading of ampicillin is hampered by

a relatively large quantity of D-phenylglycine,
relative to ampicillin, being present in the reaction
mixture obtained after the enzymatic acylation
reaction, as a result of which a smaller quantity of
5 ampicillin can be isolated.

It has been found that in order to achieve
a high conversion in the process it is of great
importance to be able to carry out the reaction at high
concentrations, and therefore also at a high
10 concentration of β -lactam nucleus.

WO-A-96/02663 describes a process in which
the enzymatic acylation reaction of β -lactam nuclei is
carried out at a constant concentration of the
reactants. In the continuous process described here the
15 aim is to achieve the highest possible level of
concentration of both reactants.

It has been found, however, that when the
preparation of ampicillin is carried out at a high
concentration of 6-APA, only a relatively low
20 conversion is achieved, compared with conversions which
could be achieved in the preparation of other β -lactam
derivatives, such as cephalixin.

The applicant has now surprisingly found
that by ensuring that the concentration of 6-APA in
25 dissolved form present in the reaction mixture is kept
relatively low, a higher conversion can be achieved
than when the concentration of dissolved 6-APA is
chosen to be as high as possible. Furthermore it is
found that the stirrability of the reaction mixture is

considerably better when the concentration of dissolved 6-APA is kept low.

In the context of the present invention "conversion" means the molar ratio of ampicillin formed to the quantity of 6-APA employed. The concentration of dissolved 6-APA is expressed as the quantity of 6-APA in moles per kg of reaction mixture; the total concentration, dissolved and undissolved, of 6-APA and ampicillin is expressed as the quantity of 6-APA plus ampicillin in moles per kg of total reaction mixture; apart from the solution, the total reaction mixture may contain a number of solid substances, for example 6-APA, ampicillin, phenylglycine and immobilized enzyme.

The molar ratio of acylation agent to 6-APA, i.e. the total quantity of added phenylglycine derivative divided by the total quantity of added 6-APA, expressed in moles, is less than 2.5. The molar ratio is preferably between 1.0 and 2.0, in particular between 1.2 and 1.8.

The enzymatic acylation reaction is preferably carried out as a batch process. If desired it is also possible to carry out the reaction continuously, with the concentration of dissolved 6-APA being controlled in line.

In the process according to the invention, the total concentration of 6-APA plus ampicillin (in dissolved and in undissolved form) in the reaction

mixture is made higher than 250 mM, preferably higher than 300 mM, and in particular higher than 350 mM.

During the preparation of ampicillin, the concentration of dissolved 6-APA is essentially kept
5 lower than 300 mM, preferably lower than 250 mM. At a higher concentration of the acylation agent a higher concentration of dissolved 6-APA can if necessary be chosen than at a lower concentration. This is because the reaction rate is higher at a higher concentration
10 of the acylation agent, which means that 6-APA is present at a high concentration in dissolved form for only a relatively short time.

The concentration of 6-APA present in the reaction mixture in dissolved form can be kept low in
15 various ways. One possibility of keeping the concentration of dissolved 6-APA low is to initially charge only part of the total quantity of 6-APA and add the rest during the reaction. A disadvantage of this, however, is that 6-APA then has to be added as a solid
20 - which creates practical problems. As a result, the total quantity of 6-APA is preferably initially charged in a batch process at the beginning of the reaction, after which, during the enzymatic acylation reaction, the concentration of 6-APA in the reaction mixture will
25 decrease and the concentration of ampicillin will increase. A suitable method of nevertheless achieving a low concentration of dissolved 6-APA is, for example, to keep the pH at a lower value compared with the pH at which a maximum solubility of the reactants is

achieved. A particularly suitable method of keeping the concentration of 6-APA in dissolved form low is, for example, to ensure that the concentration of the phenylglycine derivative is kept low, for example by
5 metering in the phenylglycine derivative partially in the course of the reaction.

It has in fact been found that when the phenylglycine derivative concentration is kept low, little 6-APA goes into solution, so that the
10 concentration of 6-APA in solution can be controlled by metering in the phenylglycine derivative.

Phenylglycine in activated form, for example an amide or an ester, in particular a methyl ester, can be used as the acylation agent in the
15 (enzymatic) acylation reaction. D-phenylglycine amide (PGA) is preferably used.

A particularly suitable embodiment is obtained when PGA is added in the form of a salt thereof, preferably the salt of PGA and a mineral acid,
20 for example PGA.HCl , $\text{PGA.1/2H}_2\text{SO}_4$ and PGA.HNO_3 . In this way it is in fact possible in a simple way to achieve optimum metering of the PGA by keeping the pH constant. $\text{PGA.1/2H}_2\text{SO}_4$ is preferably used, because this salt has a very high solubility.

25 The temperature at which the enzymatic acylation reaction is carried out is generally lower than 40°C , preferably between -5 and 35°C . The pH at which the enzymatic acylation reaction is carried out

is generally between 5.5 and 8.0, preferably between 6.0 and 6.8.

Any enzyme that is suitable as a catalyst in the linking reaction can in principle be used as the enzyme. Such enzymes are for example the enzymes which are known under the general name penicillin amidase or penicillin acylase. Such enzymes are described in for example J.G. Shewale et al., Process Biochemistry, August 1989, pp. 146-154, and in J.G. Shewale et al., Process Biochemistry International, June 1990, pp. 97-103. Examples of suitable enzymes are enzymes derived from Acetobacter, in particular Acetobacter pasteurianum, Aeromonas, Alcaligenes, in particular Alcaligenes faecalis, Aphanocladium, Bacillus sp., in particular Bacillus megaterium, Cephalosporium, Escherichia, in particular Escherichia coli, Flavobacterium, Fusarium, in particular Fusarium oxysporum and Fusarium solani, Kluyvera, Mycoplasma, Protaminobacter, Proteus, in particular Proteus rettgeri, Pseudomonas and Xanthomonas, in particular Xanthomonas citrii.

An immobilized enzyme is preferably used since the enzyme can then be simply separated off and re-used. A suitable immobilization technology is described in for example EP-A-222462. Another suitable technology involves immobilizing the Penicillin G acylase on a carrier which contains a gelling agent, for example gelatin, and a polymer with free amino groups, for example alginate amine, chitosan or

polyethylenimine. In addition, enzymes in crystalline form (CLEC'sTM) can also be used.

Of the immobilized enzymes which are commercially available, those which were found to be particularly suitable were, for example, the Escherichia coli enzyme from Boehringer Mannheim GmbH which is commercially available under the name Enzygel[®], the immobilized Penicillin-G acylase from Recordati and the immobilized Penicillin-G acylase from Pharma Biotechnology, Hannover.

The (enzymatic) acylation reaction and the further upgrading of the reaction mixture are in practice usually carried out in water. If desired, the reaction mixture can also contain an organic solvent or a mixture of organic solvents, preferably less than 30 vol%. Examples of organic solvents which can be used are alcohols with 1-7 C atoms, for example a monoalcohol, in particular methanol or ethanol; a diol, in particular ethyleneglycol; or a triol, in particular glycerol.

The reaction is preferably almost completely stopped when near to maximum conversion has been achieved. A suitable embodiment for achieving this is to lower the pH, preferably to a value between 4.0 and 6.3, in particular between 5.0 and 5.7. Another suitable embodiment is to lower the temperature of the reaction mixture as soon as maximum conversion is achieved. A combination of the two embodiments is also possible.

After the reaction has been almost stopped on achieving maximum conversion, the reaction mixture is usually present in the form of a suspension which contains several solid substances, for example
5 ampicillin, D-phenylglycine and immobilized enzyme. For the sake of process economics, the immobilized enzyme is preferably recovered. A suitable way of doing this is, for example, to filter the reaction mixture through a screen, while stirring, with the direction of
10 rotation of the agitator being preferably such that the suspension is pumped upwards in the centre of the agitator. Valuable components, for example AMPI and PG, can subsequently be recovered; for example with the aid of a pH shift. The mother liquor which remains contains
15 only a few byproducts, and can subsequently be recirculated if desired.

In the context of the present invention, the various components can be present in the reaction mixture either in free form or as salts. The stated pH
20 value always means the pH value measured with a pH electrode calibrated at room temperature.

The invention will be further explained by means of the examples, without, however, being limited thereto.

25

Abbreviations:

AMPI.3H₂O = ampicillin trihydrate
6-APA = 6-aminopenicillanic acid
PGA = D-phenylglycine amide

PG = D-phenylglycine

AssemblaseTM is an immobilized Escherichia coli penicillin acylase from E. coli ATCC 1105, as described in WO-A-97/04086. The immobilization has been carried out as described in EP-A-222462, with gelatin and chitosan being used as gelling agent and glutaraldehyde as cross-linker. The final activity of the Escherichia coli penicillin acylase is determined by the amount of enzyme which has been added to the activated globules, and amounted to 3 ASU/g of dry weight, with 1 ASU (Amoxicillin Synthesis Unit) being defined as the amount of enzyme which generates 1 g of Amoxicillin.3H₂O per hour from 6-APA and D-p-hydroxyphenylglycine methyl ester (HPGM) (at 20°C; 6.5% of 6-APA and 6.5% of HPGM).

15

Example I

Preparation of PGA.1/2H₂SO₄ solution.

301.6 g of PGA (2.00 mol) was suspended in 650 g of water at T = 5°C. 102.1 g of 96% H₂SO₄ (1.00 mol) was added dropwise over a period of 1 hour, with stirring, with the temperature being kept at T < 25°C by cooling.

Example II

Synthesis of Ampicillin

An enzyme reactor (1.5 l, diameter 11 cm), fitted with a screen bottom with a 175 μ m mesh, was
5 filled with 300 g net wet Assemblase[®].

A preparation reactor (1.2 l) was filled with 131.6 g of 6-APA (0.600 mol), 30.2 g of PGA (0.200 mol) and 400 ml of water ($T = 10^{\circ}\text{C}$). This mixture was stirred for 15 minutes at $T = 10^{\circ}\text{C}$ and then transferred
10 to the enzyme reactor at time $t = 0$ with 100 ml of water ($T = 10^{\circ}\text{C}$).

At $t = 0$ the agitator in the enzyme reactor was started. Over a period of 283 minutes 423.7 g (0.800 mol) of $\text{PGA} \cdot 1/2\text{H}_2\text{SO}_4$ solution was added at a
15 constant rate, with the temperature being kept at 10°C . The pH was about 6.3. From $t = 328$ minutes onwards the pH was kept at 6.3 by titration with 6N (aqueous) H_2SO_4 . At $t = 540$ minutes the quantity of Ampicillin was at a maximum and the pH was reduced to 5.6 by adding 6N
20 H_2SO_4 .

The enzyme reactor now contained:

575 mmol of AMPI (=96% relative to the amount of 6-APA used)

15 mmol of 6-APA

25 50 mmol of PGA

365 mmol of PG

The concentrations during the reaction are shown in Graph 1.

Comparative Experiment A

Synthesis of Ampicillin

An enzyme reactor (1.5 l, diameter 11 cm), fitted with a screen bottom with a 175 μ m mesh, was
5 filled with 300 g net wet Assemblase[®].

A preparation reactor (1.2 l) was filled with 143.2 g (0.950 mol) of PGA in 500 ml of water at 10°C. Over a period of 15 minutes 131.6 g of 6-APA (0.600 mol) was added in small portions at 10°C, with
10 cooling, while the pH was kept at 7.0 by titration with 6N (aqueous) H₂SO₄. A total of 54.5 ml of 6N H₂SO₄ was needed. The mixture was stirred for 15 minutes at T = 10°C and then transferred to the enzyme reactor at time t = 0 with 100 ml of water (T = 10°C). At t = 0 the
15 agitator in the enzyme reactor was started. The pH was kept at 7.0 by titration with 6N H₂SO₄. The temperature was kept at 10°C. At t = 160 minutes the quantity of Ampicillin was at a maximum and the pH was reduced to 5.6 by means of 6N H₂SO₄. A total of 147.6 ml of 6N H₂SO₄
20 was added to the enzyme reactor. The mixture was relatively viscous and difficult to stir.

The enzyme reactor now contained:

551 mmol of AMPI (= 92% relative to the amount of 6-APA used)
25 24 mmol of 6-APA
 50 mmol of PGA
 330 mmol of PG

The concentrations during the reaction are shown in Graph 2.

NEW SET OF CLAIMS

1. Process for the preparation of ampicillin in which 6-aminopenicillanic acid (6-APA) is subjected to an enzymatic acylation reaction with the aid of a phenylglycine derivative, with the total concentration of the 6-APA present in the reaction mixture, plus ampicillin, being greater than 250 mM, the concentration of 6-APA in solution being kept lower than 300 mM and the molar ratio of acylating agent to 6-APA employed, which molar ratio is defined as the total quantity of added phenylglycine derivative divided by the total quantity of added 6-APA, expressed in moles, being less than 2.5
2. Process according to Claim 1, in which the concentration of the 6-APA plus ampicillin present in the reaction mixture is greater than 300 mM.
3. Process according to Claim 1 or 2, in which the concentration of 6-APA in solution is kept lower than 250 mM.
4. Process according to any one of Claims 1-3, in which the molar ratio of the total acylating agent employed to 6-APA is less than 2.0.
5. Process according to any one of Claims 1-4, characterized in that the 6-APA and/or the phenylglycine derivative is metered in partially in the course of the enzymatic acylation reaction.

AMENDED SHEET

AMENDED CLAIMS

6. Process according to Claim 5, characterized in that the phenylglycine derivative is metered in as a salt of D-phenylglycine amide and an acid.
7. Process according to Claim 6, characterized in that the phenylglycine derivative is metered in the form of a solution of D-phenylglycine amide.1/2H₂SO₄ in water.
8. Process according to any one of Claims 5-7, characterized in that the metering of the phenylglycine derivative is controlled by means of pH measurement.
9. Process according to any one of Claims 1-8, characterized in that the pH of the reaction mixture is lowered as soon as near to maximum conversion is achieved.
10. Process according to any one of Claims 1-9, characterized in that the temperature of the reaction mixture is lowered as soon as near to maximum conversion is achieved.

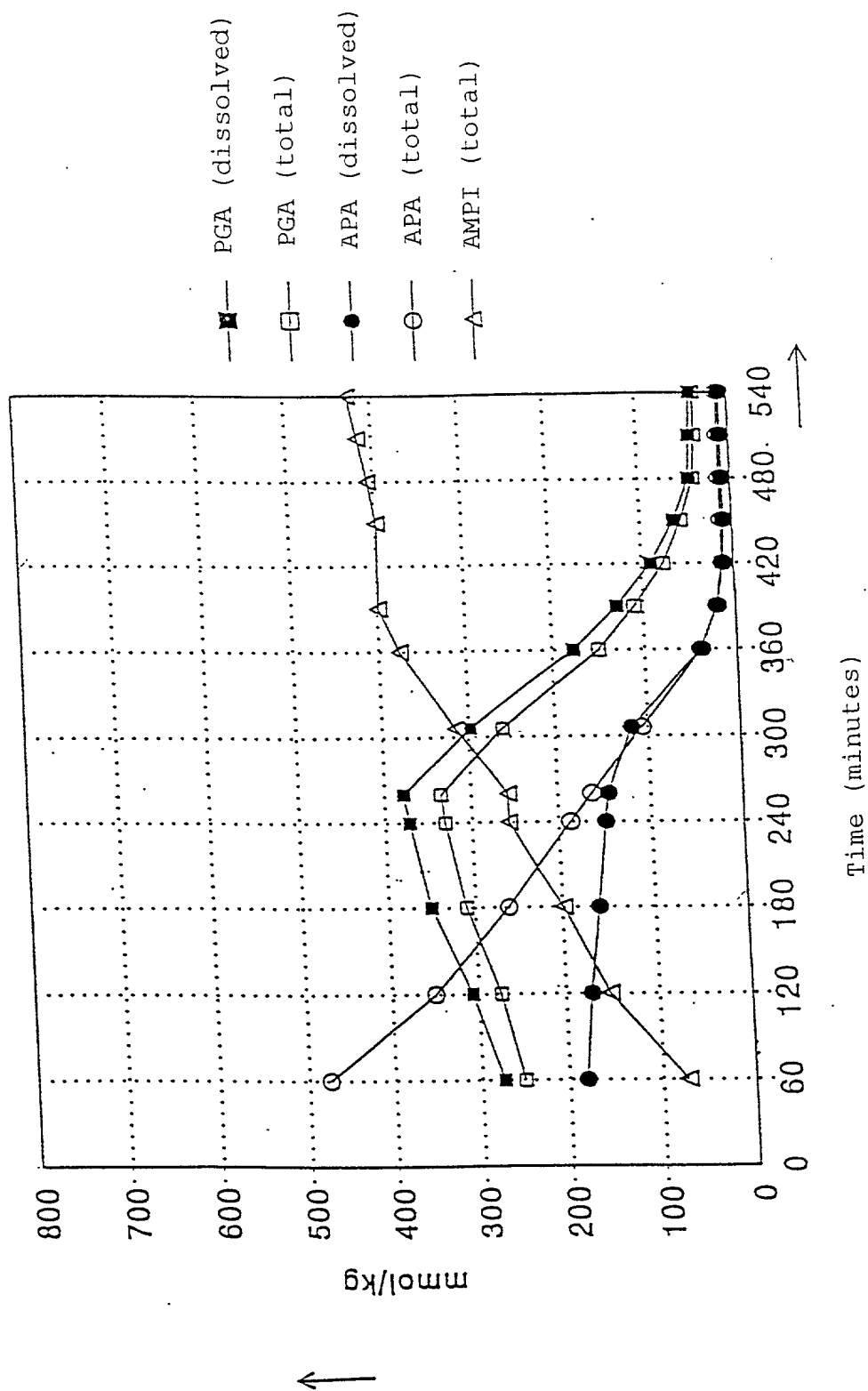
A B S T R A C T

The invention relates to a process for the preparation of ampicillin in which 6-aminopenicillanic
5 acid (6-APA) is subjected to an enzymatic acylation reaction with the aid of a phenylglycine derivative, with the total concentration of the 6-APA present in the reaction mixture, plus ampicillin, being greater than 250 mM, the concentration of 6-APA in solution
10 being kept lower than 300 mM and the molar ratio of acylation agent to 6-APA which is employed being less than 2.5.

The concentration of 6-APA present in the reaction mixture in dissolved form can be kept low in
15 various ways. One possibility of keeping the concentration of dissolved 6-APA low is to initially charge only part of the total amount of 6-APA and to add the remainder during the reaction. The total amount of 6-APA is preferably initially charged at the start
20 of the reaction. A suitable method of nevertheless achieving a low concentration of dissolved 6-APA is, for example, to keep the pH at a lower value than the pH at which maximum solubility of the reactants is achieved, by ensuring that the concentration of the
25 phenylglycine derivative is kept low, for example by metering in the phenylglycine derivative partially in the course of the reaction, for example in the form of a salt thereof, preferably the salt of PGA and a mineral acid. In this way it is possible in a simple

way to achieve optimum metering of the PGA by keeping the pH constant. $\text{PGA} \cdot 1/2\text{H}_2\text{SO}_4$ is preferably used.

GRAPH 1



GRAPH 2

